1: NM_080751. Reports Homo sapiens tran...[gi:94536851]

Links

Comment Features Sequence

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DEFINITION  Homo sapiens transmembrane channel-like 2 (TMC2), mRNA.
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VERSION     NM_080751.2  GI:94536851
KEYWORDS    .
SOURCE      Homo sapiens (human)
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            Mammalia; Eutheria; Euarchontoglires; Primates; Haplorrhini;
            Catarrhini; Hominidae; Homo.
REFERENCE   1  (bases 1 to 3169)
  AUTHORS   Kimura,K., Wakamatsu,A., Suzuki,Y., Ota,T., Nishikawa,T.,
            Yamashita,R., Yamamoto,J., Sekine,M., Tsuritani,K., Wakaguri,H.,
            Ishii,S., Sugiyama,T., Saito,K., Isono,Y., Irie,R., Kushida,N.,
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            Murakawa,K., Ishida,S., Ishibashi,T., Takahashi-Fujii,A.,
            Tanase,T., Nagai,K., Kikuchi,H., Nakai,K., Isogai,T. and Sugano,S.
  TITLE     Diversification of transcriptional modulation: large-scale
            identification and characterization of putative alternative
            promoters of human genes
  JOURNAL   Genome Res. 16 (1), 55-65 (2006)
   PUBMED   16344560
REFERENCE   2  (bases 1 to 3169)
  AUTHORS   Kurima,K., Yang,Y., Sorber,K. and Griffith,A.J.
  TITLE     Characterization of the transmembrane channel-like (TMC) gene
            family: functional clues from hearing loss and epidermodysplasia
            verruciformis
  JOURNAL   Genomics 82 (3), 300-308 (2003)
   PUBMED   12906855
REFERENCE   3  (bases 1 to 3169)
  AUTHORS   Keresztes,G., Mutai,H. and Heller,S.
  TITLE     TMC and EVER genes belong to a larger novel family, the TMC gene
            family encoding transmembrane proteins
  JOURNAL   (er) BMC Genomics 4 (1), 24 (2003)
   PUBMED   12812529
REFERENCE   4  (bases 1 to 3169)
  AUTHORS   Kurima,K., Peters,L.M., Yang,Y., Riazuddin,S., Ahmed,Z.M., Naz,S.,
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            Keats,B.J., Wilcox,E.R., Friedman,T.B. and Griffith,A.J.
  TITLE     Dominant and recessive deafness caused by mutations of a novel
            gene, TMC1, required for cochlear hair-cell function
  JOURNAL   Nat. Genet. 30 (3), 277-284 (2002)
   PUBMED   11850618
COMMENT     VALIDATED REFSEQ: This record has undergone preliminary review of
            the sequence, but has not yet been subject to final review. The
            reference sequence was derived from AF417580.2, DA769512.1 and
            AL049712.12.
            On May 4, 2006 this sequence version replaced gi:20304092.
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Summary: This gene is considered a member of a gene family predicted to encode transmembrane proteins. The specific function of this gene is unknown; however, expression in the inner ear suggests that it may be crucial for normal auditory function. Mutations in this gene may underlie hereditary disorders of balance and hearing.

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☐ 1: AF417580. Reports Homo sapiens tran...[gi:28642834]

Links

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DEFINITION  Homo sapiens transmembrane channel-like protein 2 (TMC2) mRNA,
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            Catarrhini; Hominidae; Homo.
REFERENCE   1  (bases 1 to 3169)
  AUTHORS   Kurima,K., Peters,L.M., Yang,Y., Riazuddin,S., Ahmed,Z.M., Naz,S.,
            Arnaud,D., Drury,S., Mo,J., Makishima,T., Ghosh,M., Menon,P.S.N.,
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  TITLE     Dominant and recessive deafness caused by mutations of a novel
            gene, TMC1, required for cochlear hair-cell function
  JOURNAL   Nat. Genet. 30 (3), 277-284 (2002)
   PUBMED   11850618
REFERENCE   2  (bases 1 to 3169)
  AUTHORS   Kurima,K., Griffith,A.J. and Friedman,T.B.
  TITLE     Direct Submission
  JOURNAL   Submitted (10-SEP-2001) NIDCD, NIH, 5 Research Court, #2A02,
            Rockville, MD 20850, USA
REFERENCE   3  (bases 1 to 3169)
  AUTHORS   Kurima,K., Griffith,A.J. and Friedman,T.B.
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AF417580

LOCUS AF417580 3169 bp mRNA linear PRI 05-MAR-2003
DEFINITION Homo sapiens transmembrane channel-like protein 2 (TMC2) mRNA, complete cds.
ACCESSION AF417580
VERSION AF417580.2 GI:28642834
KEYWORDS .
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae; Homo.
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AUTHORS Kurima,K., Peters,L.M., Yang,Y., Riazuddin,S., Ahmed,Z.M., Naz,S., Arnaud,D., Drury,S., Mo,J., Makishima,T., Ghosh,M., Menon,P.S.N., Deshmukh,D., Oddoux,C., Ostrer,H., Khan,S., Raizuddin,S., Deininger,P.L., Hampton,L.L., Sullivan,S.L., Battey,J.F., Keats,B.J.B., Wilcox,E.R., Friedman,T.B. and Griffith,A.J.
TITLE Dominant and recessive deafness caused by mutations of a novel gene, TMC1, required for cochlear hair-cell function
JOURNAL Nat. Genet. 30 (3), 277-284 (2002)
PUBMED 11850618
REFERENCE 2 (bases 1 to 3169)
AUTHORS Kurima,K., Griffith,A.J. and Friedman,T.B.
TITLE Direct Submission
JOURNAL Submitted (10-SEP-2001) NIDCD, NIH, 5 Research Court, #2A02, Rockville, MD 20850, USA
REFERENCE 3 (bases 1 to 3169)
AUTHORS Kurima,K., Griffith,A.J. and Friedman,T.B.
TITLE Direct Submission
JOURNAL Submitted (03-MAR-2003) NIDCD, NIH, 5 Research Court, #2A02, Rockville, MD 20850, USA
REMARK Sequence update by submitter
COMMENT On Mar 3, 2003 this sequence version replaced gi:19223982.
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ORIGIN

Query Match 100.0%; Score 3169; DB 8; Length 3169;
Best Local Similarity 100.0%; Pred. No. 0;
Matches 3169; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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The record has been replaced by NM_080751.2